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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.	
10/809,395	03/26/2004	Masaya Yamanouchi	0020-4935PUS2	8550	•
	7590 07/10/2007 ART KOLASCH & BIJ	07/10/2007 ASCH & BIRCH		EXAMINER	
PO BOX 747		*	SAUCIER, SANDRA E		
FALLS CHURCE	FALLS CHURCH, VA 22040-0747	•	ART UNIT	PAPER NUMBER	
		*	1651		
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			NOTIFICATION DATE	DELIVERY MODE	
			07/10/2007	ELECTRONIC	

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

Notice of the Office communication was sent electronically on above-indicated "Notification Date" to the following e-mail address(es):

mailroom@bskb.com

		Application No.	Applicant(s)			
		10/809,395	YAMANOUCHI ET AL.			
	Office Action Summary	Examiner	Art Unit			
		Sandra Saucier	1651			
Period fo	The MAILING DATE of this communication app or Reply	ears on the cover sheet with th	e correspondence address			
A SH WHIC - Exter after - If NC - Failu Any	ORTENED STATUTORY PERIOD FOR REPLY CHEVER IS LONGER, FROM THE MAILING DAnsions of time may be available under the provisions of 37 CFR 1.13 SIX (6) MONTHS from the mailing date of this communication. It period for reply is specified above, the maximum statutory period were to reply within the set or extended period for reply will, by statute, reply received by the Office later than three months after the mailing and patent term adjustment. See 37 CFR 1.704(b).	ATE OF THIS COMMUNICATI 36(a). In no event, however, may a reply be fill apply and will expire SIX (6) MONTHS for cause the application to become ABANDO	ON. e timely filed rom the mailing date of this communication. DNED (35 U.S.C. § 133).			
Status						
2a)⊠	Responsive to communication(s) filed on <u>25 Ar</u> This action is FINAL . 2b) This Since this application is in condition for allowar closed in accordance with the practice under E	action is non-final.				
Disnositi	on of Claims	•				
5)□ 6)⊠ 7)□	 4) Claim(s) 1-4 and 6 is/are pending in the application. 4a) Of the above claim(s) is/are withdrawn from consideration. 5) Claim(s) is/are allowed. 6) Claim(s) 1-4 and 6 is/are rejected. 7) Claim(s) is/are objected to. 8) Claim(s) are subject to restriction and/or election requirement. 					
Applicati	on Papers	,				
10)⊠	 9) The specification is objected to by the Examiner. 10) The drawing(s) filed on 25 April 2007 is/are: a) accepted or b) objected to by the Examiner. Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a). Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d). 11) The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152. 					
Priority u	ınder 35 U.S.C. § 119					
 12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f). a) All b) Some * c) None of: 1. Certified copies of the priority documents have been received. 2. Certified copies of the priority documents have been received in Application No. 09979765. 3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)). * See the attached detailed Office action for a list of the certified copies not received. 						
2) Notice 3) Information	e of References Cited (PTO-892) se of Draftsperson's Patent Drawing Review (PTO-948) mation Disclosure Statement(s) (PTO/SB/08) r No(s)/Mail Date	4) Interview Summ Paper No(s)/Mai 5) Notice of Inform: 6) Other:	I Date			

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DETAILED ACTION

Claims 1-4 and 6 are pending and are considered on the merits.

Claim Rejections - 35 USC § 103

Claims 1-4 and 6 remain/are rejected under 35 U.S.C. 103(a) as being unpatentable over US 2002/0115699 [A] in light of Mukherjee *et al.* [W2].

The claims are directed to treatment of glomerulonephritis, nephrotic syndrome, focal glomerulosclerosis, immune complex nephropathy, lupus nephritis, drug-induce renal injury and renal insufficiency, not caused by diabetes comprising administering a PPAR agonist with also has the activity of up-regulating the expression of L-FABP.

US 2002/0115699 discloses administering thiazolidinedione compounds which include isaglitazone [0002,0004,0009] to treat and prevent renal diseases including those not associated with diabetes [0005]. Although the reference is silent with regard to the PPAR agonist activity of the thiazolidinediones used to treat renal disease, this activity is inherent in this class of compounds. Thus, administering a thiazolidinedione which has known antidiabetic and renal treatment treating activity, necessarily incorporates administering PPARgamma agonist activity since these same compounds have multiple effects. Further, at least some thiazolidinediones are known to also have L-FABP up-regulating activity (Sidaway *et al.* [U]). This is an inherency rejection.

Mukherjee *et al.* disclose that thiazolidinediones in addition to their antidiabetic activity have PPARgamma agonist activity (abstract).

"To invalidate a patent by anticipation, a prior art reference normally needs to disclose each and every limitation of the claim. See Standard Havens Prods., Inc. v. Gencor Indus., Inc., 953 F.2d 1360, 1369, 21 USPQ2d 1321, 1328 (Fed. Cir. 1991). However, a prior art reference may anticipate when the claim limitation or limitations not expressly found in that reference are

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nonetheless inherent in it. See id.; Verdegaal Bros., Inc. v. Union Oil Co. of Cal., 814 F.2d 628, 630, 2 USPQ2d 1051,1053 (Fed. Cir. 1987). Under the principles of inherency, if the prior art necessarily functions in accordance with, or includes, the claimed limitations, it anticipates. See In re King, 801 F.2d 1324, 1326, 231 USPQ 136, 138 (Fed. Cir. 1986). Inherency is not necessarily coterminous with the knowledge of those of ordinary skill in the art. See Titanium Metals, 778 F.2d at 780. Artisans of ordinary skill may not recognize the inherent characteristics or functioning of the prior art. See id. at 782. However, the discovery of a previously unappreciated property of a prior art composition, or of a scientific explanation for the prior art's functioning, does not render the old composition patentably new to the discoverer. See id. at 782 ("Congress has not seen fit to permit the patenting of an old [composition], known to others . . . , by one who has discovered its . . . useful properties."); Verdegaal Bros., 814 F.2d at 633.

This court's decision in Titanium Metals illustrates these principles. See Titanium Metals, 778 F.2d at 775. In Titanium Metals, the patent applicants sought a patent for a titanium alloy containing various ranges of nickel, molybdenum, iron, and titanium. The claims also required that the alloy be "characterized by good corrosion resistance in hot brine environments." Titanium Metals, 778 F.2d at 776. A prior art reference disclosed a titanium alloy falling within the claimed ranges, but did not disclose any corrosionresistant properties. This court affirmed a decision of the PTO Board of Appeals finding the claimed invention unpatentable as anticipated. This court concluded that the claimed alloy was not novel, noting that "it is immaterial, on the issue of their novelty, what inherent properties the alloys have or whether these applicants discovered certain inherent properties." Id. at 782. This same reasoning holds true when it is not a property, but an ingredient, which is inherently contained in the prior art. The public remains free to make, use, or sell prior art compositions or processes, regardless of whether or not they understand their complete makeup or the underlying scientific principles which allow them to operate. The doctrine of anticipation by inherency, among other

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doctrines, enforces that basic principle." See Atlas Powder Co. v. IRECO Inc. 51 USPQ2d 1943 (Fed. Cir. 1999).

Thus, a reference may be anticipatory if it discloses every limitation of the claimed invention either explicitly or inherently. A reference includes an inherent characteristic if that characteristic is the "natural result" flowing from the reference's explicitly explicated limitations. Continental Can Co. USA, Inc. v. Monsanto Co., 948 F.2d 1264, 1269, 20 USPQ2d 1746, 1749 (Fed. Cir. 1991).

In the instant case, the administration of a PPAR agonist which has L-FABP up-regulating activity flows from the administration of the thiazolidinedione class of compounds, which are known to have such activity.

Response to Arguments

Applicant's arguments filed 4/25/07 have been fully considered but they are not fully persuasive.

Applicants argue that WO 2002/0115699 is directed to treatment of nephropathy caused by diabetes and that the instant claims exclude such secondary diseases caused by diabetes.

While the exemplification of WO 2002/0115699 uses Zucker rats which are 2–3 months old, which are pre-hyperglycemic at this age, see Coimbra *et al.* (V), the generic portion of the specification states at [0005] that "compounds having insulin sensitiser activity can prevent hydronephrosis and proteinuria, such as albuminuria, that that they are therefore of potential use in the treatment and/or prophylaxis of renal disease,...". This is a disclosure of the broad use in animals not necessarily pre-diabetic or diabetic. The specification goes on to say that these compounds are useful "especially renal disease associated with Type II diabetes...". However, this does not negate the fact that the specification broadly teaches the compounds' use in all types of renal disease.

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Thus applicants arguments that the reference is "mainly" concerned with the treatment of diabetic nephropathy, while it may be correct, does not negate the fact that the specification broadly teaches the favored compounds, which are thiazolidinediones, administration in all types of renal disease. The formula (I) on page 1 includes the instantly exemplified, isaglitazone. Thus, applicants' arguments that the instant claims are not directed to the treatment of the renal diseases that are a consequence of diabetes and that WO 2002/0115699 is directed to the treatment of the renal diseases that are a consequence of diabetes is not persuasive.

Any evidence that might show that there is a difference with regard to the efficacy of the different troglitazones in the treatment of glomerulonephritis or another of the renal diseases of claim 1, would be carefully considered.

Conclusion

THIS ACTION IS MADE FINAL. Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the mailing date of this final action.

Applicant should specifically point out the support for any amendments made to the disclosure, including the claims (MPEP 714.02 and 2163.06). Due to the procedure outlined in MPEP 2163.06 for interpreting claims, it is noted that other art may be applicable under 35 USC 102 or 35 USC 103(a) once the aforementioned issue(s) is/are addressed.

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Any inquiry concerning this communication or earlier communications from the examiner should be directed to Sandra Saucier whose telephone number is (571) 272-0922. The examiner can normally be reached on Monday, Tuesday, Wednesday.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, M. Wityshyn can be reached on (571) 272-0926. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

Sandra Saucier Primary Examiner

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June 28, 2007